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## Amendments to the Claims

This listing of claims will replace all prior versions, and listings, of claims in the application:

## **Listing of Claims:**

| 1.              | (Cancelled) A method for preparing a catalyst support comprising: |   |
|-----------------|---|---|
| •               | <del>(a)</del>  | combining one or more refractory oxide precursors with one or more structural   |
| <del>prom</del> | <del>oter pre</del>   | cursors to yield a precursor mixture;   |
|                 | <del>(b)</del>  | forming a support precursor from the precursor mixture; and                     |
|                 | <del>(c)</del>  | ealcining the support precursor to obtain a hydrothermally-stable structurally- |
| <del>prom</del> | <del>oted r</del> e   | efractory-oxide catalyst support, wherein the catalyst support comprises the    |
| hydro           | therma  | lly-stable structurally-promoted refractory-oxide catalyst support.             |

- 2. (Currently amended) The method\_catalyst of claim 1 15 wherein step (b) comprises hydrolysis of the one or more refractory-oxide precursors.
- 3. (Currently amended) The <u>method\_catalyst</u> of claim 1 15 wherein step (b) comprises cohydrolysis of the one or more refractory-oxide precursors and the one or more structural promoter precursors.
- 4. (Currently amended) The method\_catalyst of claim 1 15 wherein step (b) comprises precipitation of the one or more refractory-oxide precursors.

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5. (Currently amended) The method\_catalyst of claim + 15 wherein step (b) comprises coprecipitation of the one or more refractory-oxide precursors and the one or more structural

promoter precursors.

6. (Currently amended) The method-catalyst of claim 1 15 wherein the precursor mixture

comprises a sol and step (b) comprises gelling the sol.

7. (Currently amended) The method-catalyst of claim 1 15 wherein the support precursor

comprises alumina and the one or more refractory-oxide precursors comprise alumina precursors.

8. (Currently amended) The method\_catalyst of claim 7 wherein the one or more refractory-

oxide precursors are inorganic precursors, wherein the inorganic precursors comprise at least one

precursor selected from the group consisting of aluminum nitrate, aluminum sulfate, sodium

aluminate, and aluminum chloride.

9. (Currently amended) The method-catalyst of claim 7 wherein the one or more refractory-

oxide precursors comprise at least one aluminum alkoxide.

10. (Currently amended) The method-catalyst of claim 7 wherein the hydrothermally-stable

structurally-promoted refractory-oxide catalyst support is a modified alumina support.

11. (Currently amended) The method-catalyst of claim 7 wherein the precursor mixture

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comprises a sol, and wherein step (b) comprises gelling the sol.

- 12. (Currently amended) The method\_catalyst of claim 1 11 wherein gelling the sol occurs at a temperature between about 70 °C and about 100 °C.
- 13. (Currently amended) The method\_catalyst of claim 1 15 wherein the one or more structural promoter precursors comprise at least one element selected from the group consisting of W, Ta, Nb, Th, Ge, U, Sn, Sb, V, Hf, Na, K, B, Mg, Si, Ca, Ti, Cr, Mn, Fe, Co, Ni, Cu, Zn, Ga, Sr, Zr, Ba, and the lanthanides.
- 14. (Currently amended) The <u>method\_catalyst</u> of claim 1 15 wherein step (b) further comprises steaming the support precursor.
- 15. (Currently amended) A catalyst comprising a catalytic metal dispersed on the <u>a</u> catalyst support, wherein the catalyst support is prepared by the <u>a</u> method of claim 1 comprising:
- (a) combining one or more refractory-oxide precursors with one or more structural promoter precursors to yield a precursor mixture;
  - (b) forming a support precursor from the precursor mixture; and
- (c) calcining the support precursor to obtain a hydrothermally-stable structurally-promoted refractory-oxide catalyst support, wherein the catalyst support comprises the hydrothermally-stable structurally-promoted refractory-oxide catalyst support.

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- 16. (Original) The catalyst according to claim 15 wherein the catalytic metal comprises Co, Ni, Fe, Ru, or combinations thereof.
- 17. (Withdrawn) A hydrothermal reaction process comprising contacting a feed stream with the catalyst of claim 16.
- 18. (Withdrawn) The process according to claim 17 wherein the feed stream comprises synthesis gas and the catalytic metal comprises cobalt.
- 19. (Withdrawn) The process of claim 18 further comprising converting at least a portion of the synthesis gas to hydrocarbons.
- 20. (Cancelled) A method for preparing a hydrothermally-stable structurally promoted refractory exide eatalyst support comprising:
  - (a) forming a refractory-oxide material as a slurry or sol from one or more refractory oxide precursors;
- (b) adding one or more structural promoter precursors to the refractory oxide
  - (e) calcining the refractory-oxide material to obtain the hydrothermally stable structurally promoted refractory oxide eatalyst support.
- 21. (Currently amended) The method <u>Fischer-Tropsch</u> catalyst of claim 20 44 wherein step

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(a) comprises hydrolysis of the one or more refractory-oxide precursors.

22. (Currently amended) The method Fischer-Tropsch catalyst of claim 20 44 wherein step

(a) and step (b) comprise co-hydrolysis of the one or more refractory-oxide precursors and the

one or more structural promoter precursors.

23. (Currently amended) The method Fischer-Tropsch catalyst of claim 20 44 wherein step

(a) comprises precipitation of the one or more refractory-oxide precursors.

24. (Currently amended) The method Fischer-Tropsch catalyst of claim 20 44 wherein step

(a) and step (b) comprise co-precipitation of the one or more refractory-oxide precursors and the

one or more structural promoter precursors.

25. (Currently amended) The method Fischer-Tropsch catalyst of claim 20 44 wherein step

(a) further comprises gelling the refractory-oxide material by a sol-gel process.

26. (Currently amended) The method Fischer-Tropsch catalyst of claim 25 wherein the sol-

gel process is conducted at a temperature between about 70 °C and about 100 °C.

27. (Currently amended) The method Fischer-Tropsch catalyst of claim 20 44 wherein step

(b) further comprises gelling the refractory-oxide material by a sol-gel process after adding one or

more structural promoter precursors to the refractory-oxide material.

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28. (Currently amended) The method Fischer-Tropsch catalyst of claim 27, wherein the sol-

gel process is conducted at a temperature between about 70 °C and about 100 °C.

29. (Currently amended) The method Fischer-Tropsch catalyst of claim 20 44 wherein the

refractory-oxide material comprises alumina and the one or more refractory-oxide precursors

comprise an alumina precursor.

30. (Currently amended) The method Fischer-Tropsch catalyst of claim 29 wherein the one or

more refractory-oxide precursors comprise at least one precursor selected from the group

consisting of aluminum nitrate, aluminum sulfate, sodium aluminate, and aluminum chloride,

and wherein the formation of the refractory-oxide material occurs by precipitation of the one or

more refractory-oxide precursors.

31. (Currently amended) The method Fischer-Tropsch catalyst of claim 29 wherein the one or

more refractory-oxide precursors comprise aluminum alkoxides.

32. (Currently amended) The method Fischer-Tropsch catalyst of claim 29 wherein the

hydrothermally-stable structurally-promoted refractory-oxide catalyst support is a modified

transition alumina support.

33. (Currently amended) The method Fischer-Tropsch catalyst of claim 29 wherein the one or

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more structural promoter precursors comprise at least one element selected from the group consisting of W, Ta, Nb, Th, Ge, U, Sn, Sb, V, Hf, Na, K, B, Mg, Si, Ca, Ti, Cr, Mn, Fe, Co, Ni, Cu, Zn, Ga, Sr, Zr, Ba, and the lanthanides.

- 34. (Currently amended) The method <u>Fischer-Tropsch</u> catalyst of claim 29 wherein step (c) further comprises treating the refractory-oxide material to a steam treatment prior to calcination.
- 35. (Currently amended) The method <u>Fischer-Tropsch</u> catalyst of claim 29 wherein the calcination takes place at a temperature between about 400 °C and about 900°C.
- 36. (Currently amended) The method <u>Fischer-Tropsch catalyst</u> of claim 29 further comprising gelling the refractory-oxide material by a sol-gel process before or after the addition of the one or more structural promoter precursors.
- 37. (Currently amended) The method <u>Fischer-Tropsch</u> catalyst of claim 36 wherein the solgel process is conducted at a temperature between about 70 °C and about 100 °C.
- 38. (Currently amended) The method <u>Fischer-Tropsch catalyst</u> of claim 36 wherein the one or more refractory-oxide precursors comprise at least one precursor selected from the group consisting of aluminum nitrate, aluminum sulfate, sodium aluminate and aluminum chloride.
- 39. (Currently amended) The method <u>Fischer-Tropsch catalyst</u> of claim 36 wherein the one or 1856,24501 (9518,0-01)

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more refractory-oxide precursors comprise aluminum alkoxides.

- 40. (Currently amended) The <u>method Fischer-Tropsch catalyst</u> of claim 36 wherein the hydrothermally-stable structurally-promoted refractory-oxide catalyst support is a modified transition alumina support.
- 41. (Currently amended) The method <u>Fischer-Tropsch catalyst</u> of claim 36 wherein the one or more structural promoter precursors comprise at least one element selected from the group consisting of W, Ta, Nb, Th, Ge, U, Sn, Sb, V, Hf, Na, K, B, Mg, Si, Ca, Ti, Cr, Mn, Fe, Co, Ni, Cu, Zn, Ga, Sr, Zr, Ba, and the lanthanides.
- 42. (Currently amended) The method <u>Fischer-Tropsch catalyst</u> of claim 36 wherein formation of the refractory-oxide material occurs by co-precipitation of the one or more refractory-oxide precursors and the one or more structural promoter precursors.
- 43. (Currently amended) The method Fischer-Tropsch catalyst of claim 36 44 wherein the calcination takes place at a temperature of between about 400 °C and about 900 °C.
- 44. (Currently amended) A Fischer-Tropsch catalyst comprising

  a hydrothermally-stable structurally-promoted refractory-oxide catalyst support; and

  a catalytic metal effective in catalyzing a Fischer-Tropsch reaction, said catalytic metal

  being dispersed on said catalyst support.

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wherein the hydrothermally-stable structurally-promoted refractory-oxide catalyst support is prepared by a method comprising

- (a) forming a refractory-oxide material as a slurry or sol from one or more refractory-oxide precursors;
- (b) adding one or more structural promoter precursors to the refractory-oxide material; and
  - (c) calcining the refractory-oxide material to obtain the hydrothermally-stable structurally-promoted refractory-oxide catalyst support.
- 45. (Currently amended) The Fischer-Tropsch catalyst of claim 44 wherein the hydrothermally-stable structurally-promoted refractory-oxide catalyst support emprises alumina undergoes less than a 20 percent change in surface area when subjected to water partial pressures in excess of 5 bars absolute at temperatures greater than 200°C for periods of time in excess of 2 hours.
- 46. (Original) The Fischer-Tropsch catalyst of claim 44 wherein the catalytic metal comprises cobalt, nickel, ruthenium, iron, or combinations thereof.
- 47. (Original) The Fischer-Tropsch catalyst of claim 44 further comprising one or more catalytic promoters selected from the group consisting of Re, Na, K, Rb, Cs, Mg, Ca, Sr, Ba, Cu, Ag, Au Sc, Y, La, Ti, Zr, Hf, V, Nb, Ta, Pd, Rh, Os, Ir, Pt, Mn, B, Ru, P, and combinations thereof.

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- 48. (Original) The Fischer-Tropsch catalyst of claim 47 wherein the one or more catalytic promoters are selected from the group consisting of platinum, ruthenium, copper, silver, boron, and phosphorous.
- 49. (New) The Fischer-Tropsch catalyst of claim 15 wherein the calcination takes place in air at a temperature between about 450 °C and about 850°C.
- 50. (New) The Fischer-Tropsch catalyst of claim 15 wherein the hydrothermally-stable catalyst support undergoes less than a 20 percent change in surface area when subjected to water partial pressures in excess of 5 bars absolute at temperatures greater than 200°C for periods of time in excess of 2 hours.